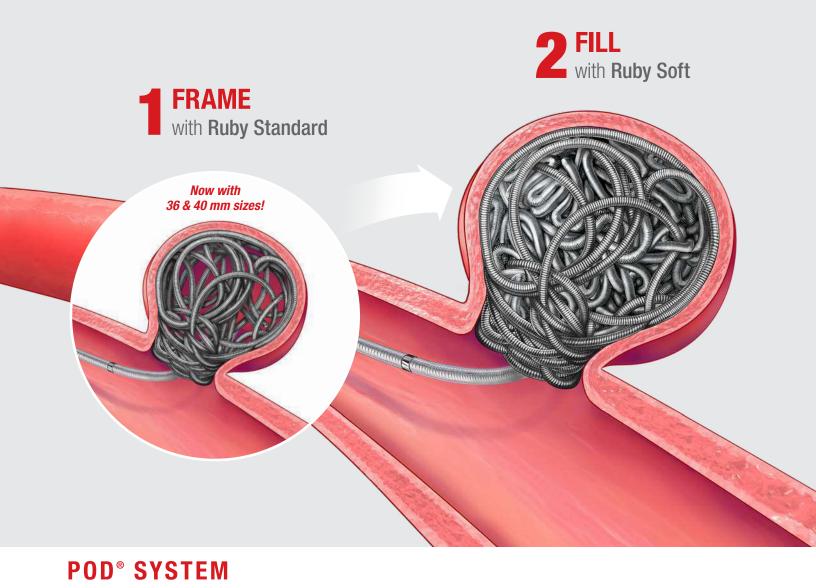


EMBOLIZATION System

FLEEDING

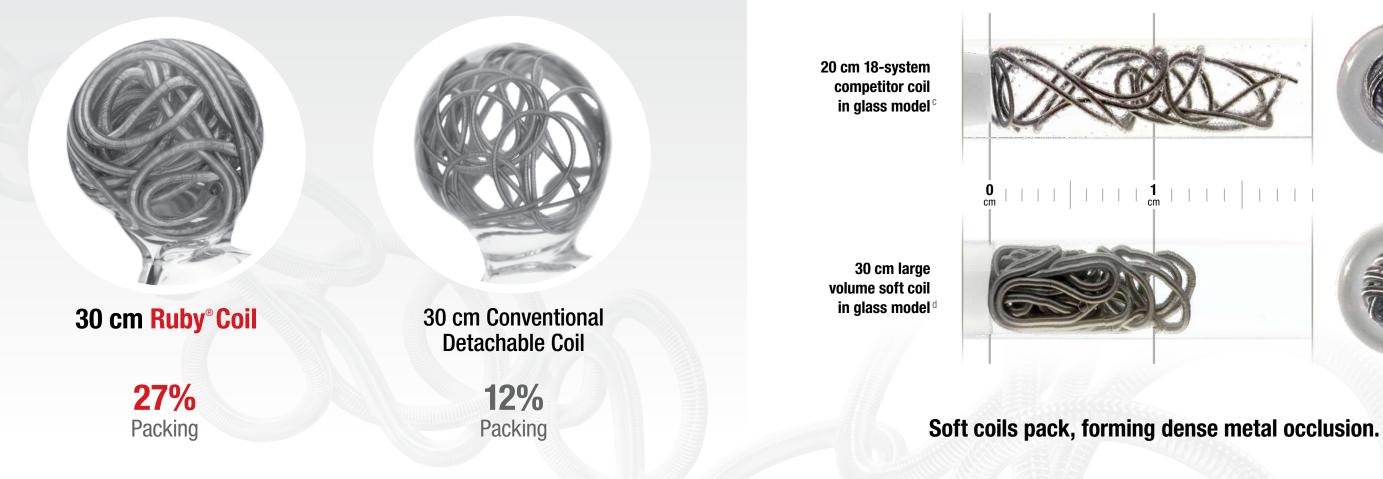
RUBY® COIL SYSTEM





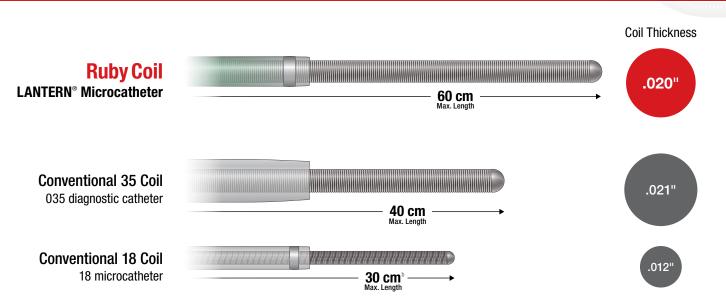
VOLUME ADVANTAGE[®]

SOFTNESS ADVANTAGE



a. 7.5 mm glass aneurysm. Tests performed and data on file at Penumbra, Inc. Bench test results may not be indicative of clinical performance.

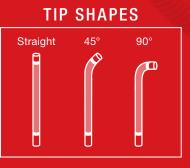
ONLY COIL TO PROVIDE 60 CM LENGTH AND .020" THICKNESS



LANTERN MICROCATHETER

c. Terumo™ AZUR® CX Coil 4 mm x 20 cm delivered through Terumo Progreat® Microcatheter into 4 mm glass tube. d. 30 cm POD Packing Coil delivered through Penumbra LANTERN® High-Flow Microcatheter into 4 mm glass tube. Photographs taken by and on file at Penumbra, Inc. Bench test results may not be indicative of clinical performance.

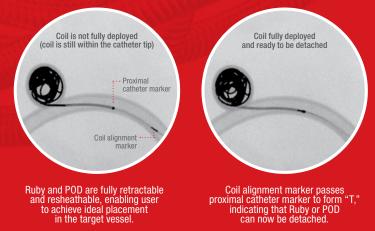
Ruby and POD[®] are the only coils designed to be delivered through a LANTERN High-Flow Microcatheter.





PRECISE COIL DETACHMENT

Ruby Coil deployed from LANTERN High-Flow Microcatheter



Photographs taken by and on file at Penumbra, Inc. Bench test results may not be indicative of clinical perform

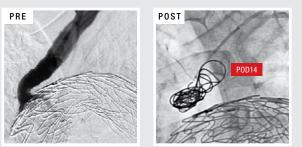
EMBOLIZATION SYSTEM CASES

Splenic Artery Embolization



Dr. Thomas Aquisto Evanston Hospital, IL

Left Subclavian Embolization



Dr. Frank Arko Atrium Health, NC

Bilateral Iliac Embolization



Dr. Herbert Cordero St. Rose Dominican Siena Campus, NV

Pelvic Congestion Syndrome



Dr. Parag J. Patel Medical College of Wisconsin, WI

Hypogastric Artery Embolization



PRE

Dr. Corey Teigen Sanford Health, ND

Hepatic Artery Aneurysm



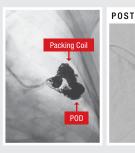
Dr. James Benenati Miami Cardiac and Vascular Institute, FL

Pre Fontan Embolization



Dr. Saar Danon Cardinal Glennon Children's Hospital, MO





Dr. Mahmood Razavi St. Joseph Hospital, CA

GDA & Gastric Embolization



Dr. Dmitri Samoilov Medical Center Radiologists, VA

ORDERING INFORMATION

RUBY COIL SYSTEM											
COMPLEX STANDARD Frame			COMPLEX SOFT Fill								
	condary iameter		Length ≀⊱	.020 in (.51 mm)							
Catalog Number	Secondary Diameter (mm)	Length (cm)	Catalog Number	Secondary Diameter (mm)	Length (cm)						
RBY2C0305 RBY2C0312 RBY2C0320	3	5 12 20	RBY4C0201 RBY4C0202 RBY4C0204	2	1 2 4						
RBY2C0410 RBY2C0420 RBY2C0435	4	10 20 35	RBY4C0305 RBY4C0315 RBY4C0406	3	5 15 6						
RBY2C0512 RBY2C0530	5	12 30	RBY4C0415 RBY4C0620	4	15 20						
RBY2C0620 RBY2C0630 RBY2C0725	6 7	20 30 25	RBY4C0630 RBY4C0835 RBY4C0860	8	30 35 60						
RBY2C0825 RBY2C0840 RBY2C1035	8	25 40 35	RBY4C1650 RBY4C2060	16 20	50 60						
RBY2C1260 RBY2C1460	12 14	60 60	DETAC	HMENT HAND)LE						
RBY2C1660 RBY2C1857 RBY2C2060	16 18 20	60 60 60			_						
RBY2C2457 RBY2C2860	24 28	60 60		Penumbra 🖗							
RBY2C3260 RBY2C3660 - Nov RBY2C4060 - Nov		60 60 60	Catalog Number RH1	Descrip Detachmen							

POD				POD PACKING COIL			
High-Flow Vessel Sacrifice				Pack Behind Ruby or POD Backstop			
Target Vessel	175 cm ∳		ength —► ⊈∫⊅		175 cm ┿ Leng	th —=	
Catalog Number		Target Vessel (mm)	Length (cm)	Catalog Number		Length (cm)	
RBYPOD3 - Now!	POD3	3	20	RBYPODJ5 - Now!	POD Packing Coil J-Soft 5 cm	5	
RBYPOD4	POD4	3.25-4	30	RBYPODJ15	POD Packing Coil J-Soft 15 cm	15	
RBYPOD5	POD5	4-5	30	RBYPODJ30	POD Packing Coil J-Soft 30 cm	30	
RBYPOD6	POD6	5-6	50	RBYPODJ45	POD Packing Coil J-Soft 45 cm	45	
RBYPOD8	POD8	6-8	60	RBYPODJ60	POD Packing Coil J-Soft 60 cm	60	
RBYPOD10 - Now!	POD10	8-10	60				
RBYPOD12 - Now!	POD12	10-12	60				
nDTFODTZ = NOW:	10012						

LANTERN Delivery Microcatheter



Indication For Use The Ruby Coil System is indicated for arterial and venous embolizations in the peripheral vasculature.

Contraindications There are no known contraindications

Warnings The Ruby Coil System should only be used by physicians who have received appropriate training in interventional techniques.

Precautions

- The device is intended for single use only. Do not resterilize or reuse. Resterilization and/or reuse may compromise the structural integrity of the device or increase the risk of contamination or infection leading. to device failure and/or cross-infection and potential
- to device failure and/or cross-infection and potential patient injury. liness or death. Do not use kinked or damaged devices. Do not use opened or damaged packages. Return all damaged devices and packaging to the manufacturer/distributor. Use prior to the "Use By" date. Use device in conjunction with fluoroscopic guidance. Do not advance or with/draw the device against resistance without careful assessment of the cause using fluoroscopy.

Indication For Use

- The POD System is indicated for the embolization of: Intracranial aneurysms
- Other neurovascular abnormalities such as arteriovenous malformations and arteriovenous fistulae
- Arterial and venous embolizations in the peripheral vasculature
- For POD Coils with nominal sizes > 6 mm The POD System is indicated for arterial and venous embolizations in the peripheral vasculature

Contraindications There are no known contraindications

- Warnings The POD System should only be used by physicians who have received appropriate training in interventional techniques.
- The device is intended for single use only. Do not resterilize or reuse. Resterilization and/or reuse may compromise the structural integrity of the device or increase the risk of contamination or infection leading to device failure and/or. cross-infection and potential patient injury, illness or death • Do not use kinked or damaged devices. Do not use

Indication For Use

media, and therapeutic devices, such as occlusion coils to the peripheral and neuro vasculature

- There are no known contraindications
- In devices are interioded for single use only. Do not resterize or ruse, Resteriziation and/or ruse may result in ineffective catheter coating lubrication, which may result in high friction and the inability to access the target location. Do not use kinked or damaged devices. Do not use open or damaged packages, Return all damaged devices and rendering the two environment of devices in the second context of the second rest of the second device. To not use open or damaged packages, Return all damaged devices and rendering the two environments of devices in the second devices of the second devices of the second device of the second devices of the second devices of the second device of the second devices of the second devices of the second device of the second devices of the second

Do not advance or withdraw the Penumbra Delivery Microcatheters against resistance without careful Microcatheters against resistance without careful assessment of the cause using fluoroscopy, if the cause cannot be determined, withdraw the device. Moving or torquing the device against resistance may result in damage to the vessel or device. • Maintain a constant infusion of an appropriate flush solution. • If flow through the device becomes restricted, do not attempt to clear the lumen by infusion. Remove and replace the device.

Potential Adverse Events

Desible complications include, but are not limited to, the following: acute occlusion, hematoma or hemorrhage at access site; death; intracranial hemorrhage; hemorrhage; infection (a access site); distal embolization; ischemia (cardiac and/or cerebral); embolus (air, foreign body, thrombus, plaque); aneurysm perforation; false aneurysm formation; neurological deficits including stroke; vessel spasm, thrombosis, dissection, perforation or rupture; air embolism; emboli.

Penumbra, Inc. USA

One Penumbra Place Alameda, CA 94502 USA

1.888.272.4606 T 1.510.748.3200 E 1 510 748 3232 order@penumbrainc.com info@penumbrainc.com

Caution: Federal (USA) law restricts these devices to sale by or on the order of a physician. Prior to use, please refer to the Instructions for Use for Ruby Coil System, POD System, and Penumbra Delivery Microcatheters for complete product indications, contraindications, warnings, precautions, potential adverse events, and detailed instructions for use. Photographs taken by and on file at Penumbra, Inc. Tests performed and data on file at Penumbra, Inc. Bench test results may not be indicative of clinical performance. Renderings for illustrative purposes only. Individual results may vary depending on a variety of patient-specific attributes. Images used with permission. Consents on file at Penumbra, Inc. Please contact your local Penumbra representative for more information.

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solution.

anaphylaxis from contrast media; aneurysm rupture; arteriovenous fistula; cagulopathy; coil hermitation into parent vessel; death; device malfunction; distal embolization; emboli; embolic stroke and other cerebral ischemic events; false aneurysm formation, hematoma or hemorrhage at access site of entry; incomplete aneurysm occlusion; intection; intima dissection; intracrianial hemorrhage; ischemia; myocardial infarction; neurological deficits including stroke; parent artery occlusion; peripheral thromboembolic events; post-embolization envertions; compared to delabemost: detapending the ceneralization syndrome; premature device detachment; recanalization; renal failure; respiratory failure; revascularization; thromboembolic episodes; vessel spasm, thrombosis, dissection, or perforation.

· Moving or torquing the device against resistance may result in damage to the vessel or device. • Maintain a constant infusion of an appropriate flush

Potential Adverse Events Possible complications include but are not limited to: acute occlusion; air embolism; allergic reaction and anaphylaxis from contrast media; aneurysm rupture;

- Use prior to the "Use By" date
- Use device in conjunction with fluoroscopic guidance.
 Do not advance or retract the device against resistance without careful assessment of the cause using fluoroscopy. If POD cannot be advanced or retracted, withdraw the device as a unit with the microcatheter
- Moving or torquing the device against resistance may result in damage to the vessel or device.
 Maintain a constant infusion of an appropriate flush solution.
- Potential Adverse Events

Potential complications include but are not limited to: acute occlusion; air embolism; allergic reaction and anaphylaxis from contrast media; coagulopathy; coil hemiation into parent vessel; death; device malfunction; distal embloization, embloi, emblois stroke maintenti distal embloization, embloi, emblois stroke and other cerebral ischemic events; false aneurysm formation; hematoma or hemorrhage at access site of entry; incomplete aneurysm occlusion; infection; infima dissection; infracranal hemorrhage; ischemia; oissection; initia/carlial nemormage; iscnemia; myccardial infaction; neurological deficits including stroke; parent artery occlusion; peripheral thromboembolic events; post-embolization syndrome; premature device detachment; recaralization; renal failure; respiratory failure; revascularization; thromboembolic episodes; vessel events; therealized in center and and and and and and and and and events; therealized in center and and and and and and and and and failure; revascularization; thromboembolic episodes; vessel spasm, thrombosis, dissection, or perforation

- Precautions

 - opened or damaged packages. Return all damaged devices and packaging to the manufacturer/distributor

The Penumbra Delivery Microcatheters are intended to assist in the delivery of diagnostic agents, such as contrast

Contraindications

- Warnings The Penumbra Delivery Microcatheters should only be used by physicians who have received appropriate training in interventional techniques.

- Precautions The devices are intended for single use only. Do not

- a clamage photos and a clamage of the manufacturer / distributor. Use prior to the "Use By" date. Use the Penumbra Delivery Microcatheters in conjunction with fluoroscopic visualization.

